

K010918 LIFEPAK 12 DEFIBRILLATOR/MONITOR SYSTEMAug 23, 2001
149 days to decisionK010918 · Product code: **MKJ** · CardiovascularSource: <https://www.510kdatabase.net/k010918/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent - ST
Submission type	Traditional
Device classification	Automated External Defibrillators (non-wearable) (MKJ)
Date received	Mar 27, 2001
Decision date	Aug 23, 2001
Days to decision	149 days
Third-party review	No

APPLICANT

Company	Medtronic Physio-Control Corp.
Location	Redmond, WA, US
Contact	SHERRI L POCOCK
510(k) history	8 submissions · 6 cleared · 2001-2004

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k010918/>; Data sourced from FDA 510(k) public records (accessdata.fda.gov).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated May 14, 2026